EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L5	1	("5,384,250").PN.	US-PGPUB; USPAT	OR	OFF	2006/09/15 14:29
S1	20	US-4692332-\$.DID. OR US-4873316-\$.DID. OR US-4877610-\$.DID. OR US-4966753-\$.DID. OR US-4970071-\$.DID. OR US-5130415-\$.DID. OR US-5206153-\$.DID. OR US-5302698-\$.DID. OR US-5322775-\$.DID. OR US-5384250-\$.DID. OR US-5589604-\$.DID. OR US-5633076-\$.DID. OR US-5648243-\$.DID. OR US-5723585-\$.DID. OR US-5723585-\$.DID. OR US-5723585-\$.DID. OR US-5723585-\$.DID. OR US-5827690-\$.DID. OR US-5827690-\$.DID. OR US-5831141-\$.DID. OR US-5843776-\$.DID. OR	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:50
S2	4	US-5965528-\$.DID. OR US-6013857-\$.DID. OR US-6288034-\$.DID. OR US-6331611-\$.DID.	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:42
S3	10348	Alphafetoprotein or AFP or foetoprotein or fetoprotein	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:46
S4	70	S3 near8 (mutant or mutation or mutate)	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:46
S5	2	S4 with "233"	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:45
S6	2	S4 with glutamine	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:45
S7	0	S6 not S5	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:45
S8	26	Alphafoetoprotein	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:46
S9	0	S8 near8 (mutant or mutation or mutate)	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:46
S10	10348	Alphafetoprotein or AFP or foetoprotein or fetoprotein	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:50
S11	26	Alphafoetoprotein	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:50
S12	329	(S10 or S11) with fragment	US-PGPUB; USPAT	ADJ	ON	2006/09/15 12:50

EAST Search History

S13	15	S12 with domain	US-PGPUB;	ADJ	ON	2006/09/15 14:29
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9/15/06 2:36:39 PM C:\Documents and Settings\RWax\My Documents\EAST\Workspaces\10624380.wsp

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189 FETOPROTEIN

175 ALPHA(W) FETOPROTEIN

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=> index biosci

FILE 'DRUGMONOG' ACCESS NOT AUTHORIZED COST IN U.S. DOLLARS FULL ESTIMATED COST

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DESDOUETS C (Reprint); FABRE M; GAUTHIER F; BRECHOT C; SOBCZAKTHEPOT J FAC MED NECKER ENFANTS MALAD, INSERM, U370, F-75730 PARIS 15, FRANCE; HOP BICETRE, SERV CENT ANAT PATHOL, LE KREMLIN BICETR, FRANCE The Genuine Article (R) Number: TC362 PROLIFERATION AND DIFFERENTIATION OF A HUMAN HEPATOBLASTOMA TRANSPLANTED ANSWER 1 OF 2 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation a> dup rem 13 PROCESSING COMPLETED FOR L3 14
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JOURNAL OF HEPATOLOGY, (NOV 1995) Vol. 23, No. 5, pp. 569-577. CYA SO

MUNKSGAARD INT PUBL LID, 35 NORRE SOCADE, PO BOX 2148, DK-1016 COPENHAGEN DENMARK. ЬB

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* Entered STN: 1995 Last Updated on STN: 1995 Reference Count: 36 Article; Journal LIFE; CLIN English PT FS LEA REC

athymic Nude mice to provide a model system to study proliferation acid differentiation of these tumoral cells. The first transplantation selected the embryonal component of this tumor, while subsequent passages selected in addition neuroendocrine and mesenchymal cells that evolved into osteoid and bony trabeculae, The embryonal character of this hepstoblastoma was further demostrated by the expression of "**glutamine*** synthetase mRNA and a fetal pattern of mRNAs encoding insulinlike growth factor II, A pure epithelial human hepatoblastoma was directly transplanted to æ

However, alphafetoprotein mRNA was detectable in neither the original nor the transplanted tumors. Finally, although p53 mRNA levels were increased, no ""mutation"" was detected in the p53 gene. (C) Journal of Hepatology. ANSWER 2 OF 2 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation

1992:78872 SCISEARCH << LOGINID::20060915>>

The Genuine Article (R) Number: HB111 NUCLEOTIDE-SEQUENCE OF PROTHROMBIN GENE IN ABNORMAL PROTHROMBIN-PRODUCING

HEPATOCELLULAR-CARCINOMA CELL-LINES TAGAWA M; OMATA M (Reprint); OHTO M

CHIBA UNIV, SCH MED, DEPT MED 1, 1-8-1 INOHANA, CHIBA 280, "APAN (Reprint)

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CANCER, (1 FEB 1992) Vol. by, Mo. J, Pr. 1.5SN: 0008-543X.
ISSN: 0008-543X.
WILEY-LISS, DIV JOHN WILEY & SONS INC 605 THIRD AVE, NEW YORK, NY ЬВ

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Article; Journal

English

Reference Count: 36 Entered STN: 1994 E K F S E

Last Updated on STN: 1994

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

not completely gamma-carboxylated and are functionally inactive. This protein can be detected in the plasma of patients with heparocallular carcinoma (HCC) and used as a new tumor marker. To analyze the mechanism of PIVKA-II production in HCC tissue, the prothrombin gene of PIVKA-II secreting HCC call lines was sequenced to detect the "...mutetion". In the Gia domain and carboxylase recognition site of leader sequence located on exons I and II that may cause the inhibition of A protein induced by vitamin K absence or antagonist II, PIVKA-II is synthesized in the liver and possesses a structure similar to prothrombin except that ten glutamic acid residues in amino-terminal Gla domain are AB

carboxylation. Exons I and II and donor and acceptor site of introl of the prothrombin gene in two HCC cell lines, PLC/PRF/S and hiH-2, were analyzed by polymerase chain reaction (PCR), and the product was sequenced directly. In addition, RNA samples of these cell lines were used for complementary DNA synthesis, followed by PCR and sequencing. The moutdootide sequences of the Gla domain in both HCC cell lines were conserved. One nucleotide change was detected at nt.554 (admine to quanine), but this did not influence the amino acid sequence. Splicing prothrombin, and protease target sites also were conserved as the reported prothrombin gene, and mutations reported for other des-gamma-carboxy coequiation factors were not detected. These results also were confirmed by DNA analysis of seven human fresh-frozen samples (three PIVKA-II-positive HCC samples and four control specimens). The mechanism of PIVKA-II production in HCC is still unclear, but it is not caused by sites between exons I and II, the leader sequence of the precursor

in the prothrombin gene.

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FILE 'REGISTRY' ENTERED AT 13:37:52 ON 15 SEP 2006 175 S ALPHA(W) FETOPROTEIN

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3 **5** E

The Genuine Article (R) Number: 055RC Differential promoter usage for insulin-like growth factor-II gene in Chinese hepatocellular carcinoma with hepatitis B virus infection Reference Count: 49

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Author Keywords: hepatocellular carcinoma; insulin-like growth factor-ll; genes; promoters; hepatitis B virus; prognosis; p53 "**mutation*"; genes; promoters; hepatitis B virus; prognosis; p53 "**mutation"; ; hepaticin; adjacent tumor tissue; hepatocepsular invasion; "**alpha*** - "*fetoprodin***; microsatelite formation Keywords Plus (R): GAIN-OF-FUNCTION; IGF-II; MESSENGER-RNA; TRANSGENIC MICE; FUNCTION MUTATIONS; EXPRESSION; P53; HEPATOCARCINOGENESIS; TRANSCRIPTION; ACTIVATION STP

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2006:582594 SCISEARCH

The Genuine Article (R) Number: 050YB Evaluation of serum levels of p53 in hepatocellular carcinoma in Egypt

- ***fetoprotein***; hepatitis B virus Evaluation or Reference Count: 16
MEDICAL LABORATORY TECHNOLOGY AN GA TI REC CC ST

WO 2004018995 AZ WO 2003-USZ6023 20030820; US 2004053392 AI Provisional US 2002-405494P 20020823, US 2003-419462 20030421; AU 2003262727 AI AU 2003-265727 20030820; US 2005065324 AI Provisional US 2002-465494P 20020823, CIP of US 2003-419462 20030421, US 2004-782968 20040220; EP 1572225 AZ EP 2003-793149 20030820, WO 2003-USZ6023 20030820; AU 2003-262727 A8 AU 2003-262727 20030820 ő ö DE DK KP KR PG PH UZ VC New purified thrombospondin fragment extracted from a body fluid, useful for diagnosing cancer e.g. adenoma, adenocarcinoma, carcinona, lymphoma leukemia or as calibrators, indicators, immunogens and analytes. 3 ≥ ***fetoprotein*** ; Pichia pastoris; (c) 2006 The Thomson Corporation IT KE נו גד נט cloning; expression KeyWords Plus (R): YEAST; PROTEIN; DNA; CONSERVATION; ALBUMIN; REGION *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* CZ WC CZ p53 GR HU UG ZIA CO GR I S JP TZ UA HU IE IT STN ***alpha*** (HBV); hepatitis C virus (HCV); hepatocellular carcinoma; KeyWords Plus (R): HEPATITIS-C; B-VIRUS; PROGNOSIS; GENE *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORWATS* => s 18 and (nonglycosylat? or (non(w) glycolylat?))
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YR NL PT RO SE SI SK TR

YR AB BG CH CY CZ DE DK EE ES FI P

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YR AB 200509103 (2006677)

Y AZ WO 2003-US7 Genuine Article (R) Number: 035PH cloning and expression of ***human*** COPYRIGHT BIOTECHNOLOGY & APPLIED MICROBIOLOGY Author Keywords: ***alpha*** - ** Subcloning and expression of 1 L8 AND PROKARYOT? SCISEARCH SCISEARCH ANSWER 1 OF 1 WPIDS ***fetoprotein*** Reference Count: 22 2004-226901 [21] and prokaryot? ANSWER 3 OF 166 2006:417537 C2004-089523 ą a> d 111 bib B04 D16 18 6 6 REC CC ST :: STP 112 N A DC PA CYC ADT <u>1</u>.8 3 8 F

- ş 6 AU 2003262727 Al Based on WO 2004018995; EP 1572225 A2 Based 2004018995; AU 2003262727 A8 Based on WO 2004018995 US 2003-419462 20030421; US 2002-405494P 20020823 WO2004018995 A UPAB: 20040326 PRAI FDT
- NOVELTY A purified thrombospondin fragment that has been extracted from a bodily fluid, where the fragment is within a molecular weight range selected from 80-10 kDa, 40-60 kDa or 20-35 kDa, and where the size in kDa is determined by gel electrophoresis after disulfide bond reduction, is

DETAILED DESCRIPTION - A thrombospondin fragment or its portion comprising:

- residues N-230 and (a) one that starts between amino acyl residues inclusive and ends between amino acyl residues V-400
- (b) one that starts between amino acyl residues N-230 and G-253, inclusive and ends between amino acyl residues D-527 and S-551;
- (c) one that starts between amino acyl residues N-230 and G-253, inclusive and ends between amino acyl residues G-787 and V-811; (d) one that starts between amino acyl residues I-165 and V-263,
- inclusive and ends between amino acyl residues K-412 and 1-530, (e) one that starts between amino acyl residues 1-165 and V-263, inclusive, and ends between amino acyl residues 1-530 and R-733;
- (g) one that starts between amino acyl residues 1-165 and V-263, inclusive, and ends between amino acyl residues K-412 and 1-530; (f) one that starts between amino acyl residues 1-165 and V-263, inclusive, and ends between amino acyl residues R-733 and Y-982;
- V-263, (h) one that starts between amino acyl residues 1-165 and
- The thrombospondin fragment comprises at least 4-6 contiguous amino V-263, inclusive, and ends between amino acyl residues R-792 and Y-982. inclusive, and ends between amino acyl residues 1-530 and R-733; (i) one that starts between amino acyl residues 1-165 and V

acyl residues from the thrombospondin sequence, where the amino acid sequence of the fragment is limited to one that is outside of a thrombospondin region given above.

INDEPENDENT CLAIMS are also included for:

- (1) a molecule identical in primary structure to the compound above; (2) a method to detect and/or quantify a thrombospondin fragment; (3) a method of producing antibodies against a thrombospondin fragment comprising administering the fragment to an organism capable of
 - (4) a monoclonal or polyclonal antibody produced by the method of producing antibodies;
- (5) a cell line producing the monoclonal antibodies or the binding (3);
 - (6) a method of producing a peptide or non-peptide binding agent against a thrombospondin fragment; agent;
- (7) a kit for the determination of the presence of, and/or the amount of, and/or the concentration of, a thrombospondin fragment in a material taken or gathered from an organism comprising the thrombospondin fragment, a binding agent that will react with thrombospondin but not with the thrombospondin fragments of interest but not with thrombospondin; fragment or fragments of interest or an antibody that will react
 - (8) a method comprising determining the amount of the unlabeled or differently labeled fragment through comparison to the results obtained
 - from the unlabeled or differently labeled fragment; (9) a method to detect the presence and/or clinical course of
- neoplastic disease in an individual; and (10) a method of producing a binding agent against a thrombospondin

fragment comprising binding a phage to the thrombospondin fragment.

USE - The thrombospondin fragments are useful in diagnostic methods for cancer, as method calibrators, method indicators, as immunogens and as manyves for methods with sustained clinical utility. Cancer is selected from adenoma, adenocarcinoma, carcinoma, lymphoma, leukemia, solid cancer, iquid cancer, metastatic cancer, pre-metastatic cancer, non-metastatic cancer, acancer, ath vascular invasion, internal cancer, skin cancer of the respiratory system, cancer of the circulatory system, cancer of the musculoskeletal system, cancer of a muscle, cancer of a bone, cancer of a joint, cancer of a tendon or ligament, cancer of a bone, system, cancer of the liver or billary system, cancer of the pancreas, cancer of the head, cancer of the neck, cancer of the pancreas, cancer of the reproductive system, cancer of the male reproductive system, cancer of the female reproductive system, cancer of the genitourinary system, cancer of a kidney, cancer of the urinary tract, canter of a sensory system, cancer of the nervous system, cancer of a lymphoid organ, prostate gland, cancer of an endometrial tissue, cancer of a mesodermal tissue, cancer of an ectodermal tissue, cancer of an endodermal tissue, teratoma, a poorly-differentiated cancer, a well-differentiated cancer, and a moderately differentiated cancer. blood cancer, cancer of a gland, cancer of a mammary gland, cancer of a

■> d 112 bib ab

New bifunctional protein comprising polypeptide domains with desired bloactivity and encodes "**human"* "**alpha*** - ***fetoprotein*** or its fragment, useful for treating e.g., viral infection, cancer rheumatoid arthritis and rhinitis. (MEAD-I) MEADE H; (GENZ) GTC BIOTHERAPEUTICS INC ANSWER 1 OF 1 WPIDS COPYRIGHT 2006 THE THOMSON CORP on STN 2005-223393 [23] WPIDS <<LOGINID::20060915>> COX, G; MEADE, H; COX, G F (COXG-I) COX G F; (MEAD-I) 109 C2005-071646 L12 AN DNC TI DC IN CYC

WW 2005024044 A2 20050317 (200523) + EN 127

RW: AT BE BG BW CH CY CZ DE DK EA EE ES FI FR GB CH GM GR HU IE IT KE

LS 1U MC MW MZ NA N LO AP L PT RO SD SE SI SK SL SZ TR TZ UG ZM ZW

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KP RR KZ LC LK LR LS LT LU LV MA MD MG MK MN MM MK KZ NA NI NO NZ

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US 2006105347 A1 20060518 (200634)

EP 1670931 A2 20060621 (200643) EN

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PT RO SE SI SK TR

NOVELTY - A bifunctional protein encoded by a transgene ΩNA construct comprising a first polypeptide domain which has a desired bloactivity and US 2003-500910P 20030905; US 2004-933854 WO2005024044 A UPAB: 20050411 FOT PRAI

20040903

WO 2005024044 A2 WO 2004-US29128 20040903; US 2006105347 A1 Provisional US 2003-500910P 20030905, US 2004-933854 20040903; EP 1670931 A2 EP 2004-783399 20040903, WO 2004-US29128 20040903

ADT

EP 1670931 A2 Based on WO 2005024044

a second polypeptide domain which has a desired bioactivity, where the encoded polypeptide of the first polypeptide domain is ""human""
""alpha"" - ""fetoprotein"" or its fragment having the biological activity of ""human". ""alpha". ""fetoprotein". "Is new.
DETAILED DESCRIPTION - A bifunctional protein encoded by a transgene
DNA construct comprising a first polypeptide domain which has a desired
bloactivity and a second polypeptide domain which has a desired
bloactivity, rogether comprising the fusion protein where each first and
second polypeptide domains retain their desired bloactivity where the ***human***

alpha... - ...fetoprotein... or its fragment having the biological ity of ...human... ...alpha... - ...fetoprotein... is new. INDEPENDENT CLAIMS are also included for the following: encoded polypeptide of the first polypeptide domain is
""alpha"" - ""fetoprotein"" or its fragment ha
activity of ""human"" ""alpha"" - ""fetopro

(1) an isolated polynucleotide encoding the fully defined 591 amino acid (SEQ ID NO: 4) sequence given in the specification linked to any of 16 fully defined 93-464 amino acid (SEQ ID NO: 10-25) sequence given in the specification;

the fusion protein, where the vector is an expression vector comprising a promoter operably linked to the fusion protein; (2) a recombinant DNA vector comprising the nucleic acid sequence of (3) a host cell transformed with the recombinant DNA vector; (4) a fusion protein produced by a method comprising expressing the fusion protein above by a cell or transgenic animal and recovering the

the

(5) a method for the production of transgenic animals capable of protein;

producing a fusion protein of interest;
(6) the resultant offspring of the method above;
(7) the resultant milk derived from the offspring; and

(8) an isolated nucleic acid sequence comprising fully defined 2029 bp (SEQ ID NO: 1) sequence given in the specification. ACTIVITY - Virucide; Anti-HIV; Cytostatic; Antirheumatic;

Antiarthritic; Neuroprotective; Osteopathic; Antipsoriatic; Muscular-Gen.; Immunosuppersaive; Dermaclogical; Antiallergic; Antialiammatory; Vulnerary; Antidiabetic; Hepatotropic. No biological data given. MECHANISM OF ACTIO: Protein therapy.

erythematogus, Hairy cell leukemia, chronic myelogenous leukemia, cutaneous T cell lymphoma carcinoid tumors, renal cell carcinoma, squamous epithelial tumors of the head and neck, multiple myeloma, malignant melanoma, Hepatitis B, Hepatitis C or low grade non-Hodgkin lymphoma USE - The fusion protein or a composition of matter containing the fusion protein is useful for treating a disease or condition such as viral infection, preferably caused by HIV, cancer (e.g., ***human*** cancer skin disease or injury such or tumor cell), rheumatoid arthritis, multiple sclerosis, osteoporosis, psoriasis, myasthenia gravis, ***human*** skin disease or injury su as photoaging damage, rhinitis, sumburn, dermatitis and burns, muscular dystrophy, insulin-dependent diabetes mellitus, systemic lupus

ENTRY 127.48 SINCE FILE => log h COST IN U.S. DOLLARS

FULL ESTIMATED COST

SESSION 184.73

SESSION WILL BE HELD FOR 60 MINUTES STN INTERNATIONAL SESSION SUSPENDED AT 13:55:54 ON 15 SEP 2006

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: ssspta1653raw

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FILE 'SCISEARCH, BIOTECHNO, CAPLUS, EMBASE, IFIPAT, MEDLINE, TOXCENTER' ENTERED AT 13:40:19 ON 15 SEP 2006

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INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, ANTE, AQUALINE, AQUASCI, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CAPLUS, CEABA-VTB, CIN, CONFSCI, CROPES, CROPE, DDFU, DEPU, DEPU, DGENE, DISSABS, DRUGB, DRUGMONGZ, DRUGY, EMBASE, ... 'ENTERD AT 13:42:03 ON 15 SEP 2006 SEA LI AND (MUTANI OR MUTATION OR MUTATE)

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